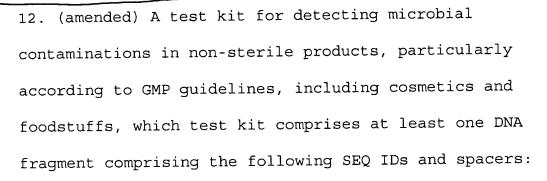
Amended Claims - Clean Copy



- a) a forward primer (SEQ ID forward primer);
- b) a probe (SEQ ID probe);
- c) a reverse primer (SEQ ID reverse primer);
- d) optionally a spacer between forward primer and probe,
- e) optionally a spacer between probe and reverse primer;
- f) optionally a spacer upstream from the forward primer,
- g) optionally a spacer downstream from the reverse primer, the SEQ Ids ((SEQ ID forward primer), (SEQ ID probe), and (SEQ ID reverse primer)) also comprising variants wherein one, two or three nucleotides have been substituted, deleted and/or inserted, the variant essentially having the same function as the sequence of the SEQ IDs ((SEQ ID forward primer), (SEQ ID probe), and (SEQ ID reverse primer)), with probes, the function of binding to DNA, and with primers, the function of binding to DNA and providing an

extendable 3' end for the DNA polymerase,

the spacers comprising 0-40 nucleotides, the DNA fragment, selected from the group of

- (i) for Pseudomonas aeruginosa

 SEQ ID No. 9 as forward primer

 SEQ ID No. 10 as probe, and

 SEQ ID No. 11 as reverse primer
- (ii) for Escherichia coli
 SEQ ID No. 12 as forward primer
 SEQ ID No. 13 as probe, and
 SEQ ID No. 14 as reverse primer
- (iii) for Salmonella ssp.
 SEQ ID No. 15 as forward primer
 SEQ ID No. 16 as probe, and
 SEQ ID No. 17 as reverse primer
- (iv) for bacteria

 SEQ ID No. 18 as forward primer

 SEQ ID No. 19 as probe, and

 reverse primer chosen from the group consisting

 of SEQ ID No. 20 and SEQ ID No. 55
- (v) for enterobacteriaceae

 SEQ ID No. 44 as forward primer

 SEQ ID No. 46 as probe, and

 SEQ ID No. 45 as reverse primer
- (vi) for enterobacteriaceae (16S rRNA)

SEQ ID No. 47 as forward primer

SEQ ID No. 48 as probe, and

SEQ ID No. 49 as reverse primer

or additionally all those sequences which are complementary to the above sequences from SEQ ID No. 9 to 49 and SEQ ID No. 55.

13. (amended) A method of detecting microorganisms in products, particularly in drugs or cosmetics, said method comprising the following steps:

 a) use of primers and fluorescence-labeled probes having the appropriate sequences and variations thereof,

(i) for Pseudomonas aeruginosa

SEQ ID No. 9 as forward primer

SEQ ID No. 10 as probe, and

SEQ ID No. 11 as reverse primer

(ii) for Escherichia coli

SEQ ID No. 12 as forward primer

SEQ ID No. 13 as probe, and

SEQ ID No. 14 as reverse primer

(iii) for Salmonella ssp.
SEQ ID No. 15 as forward primer
SEQ ID No. 16 as probe, and
SEQ ID No. 17 as reverse primer



(iv) for bacteria

SEQ ID No. 18 as forward primer

SEQ ID No. 19 as probe, and

reverse primer chosen from the group consisting

of SEQ ID No. 20 and SEQ ID No. 55

(v) for enterobacteriaceae

SEQ ID No. 44 as forward primer

SEQ ID No. 46 as probe, and

SEQ ID No. 45 as reverse primer

(vi) for enterobacteriaceae (16S rRNA)

SEQ ID No. 47 as forward primer

SEQ ID No. 48 as probe, and

SEQ ID No. 49 as reverse primer

or additionally all those sequences which are complementary to the above sequences from SEQ ID No. 9 to 49 and SEQ ID No. 55;

- b) propagating the DNA using PCR, and
- c) irradiating with specific wavelengths exciting the fluorescent dye,
- d) measuring and quantifying the emission of the excited fluorescent dye.